forming multilamellar liposome products having an average diameter of less than about 1000 nm.

- 16. The method of claim 15 wherein the multilamellar liposome products are formed by carrying out a lyophilization step.
- 17. The method of claim 15 wherein the liposomes obtained in step (b) have an everage diameter of less than about 300 nm.
- 18. The method according to claim 17 wherein the liposomes are obtained in step (b) by extrusion.
- 19. The method according to claim 15 wherein the multilamellar liposome products have an average diameter of less than about 800 nm.
- 20. The method according to claim 15 wherein the multilamellar liposome products have an average diameter of less than about 300 nm.
- 21. The method according to any one of claims 15 through 20 wherein the water soluble polymer is PEG.
- 22. The method of claim 15 wherein the amphipathic compound in a biologically active conformation is characterized as having one or more α -or π -helical domains.
- 23. The method of claim 15 wherein the biologically active amphipathic compound is a member of the vasoactive intestinal peptide (VIP)/growth hormone releasing factor (GRF) family of peptides.
 - 24. The method of claim 15 wherein the peptide is VIP.
- 25. An echogenic liposome diagnostic product manufactured by the method according to any one of claims 15 through 24.
 - 26. A diagnostic method comprising the steps of:

preparing a multila mellar liposome product comprising a biologically active amphipathic compound in association with a liposome according to the method of claim 15 through 24;

administering a diagnostically effective amount of said multilamellar liposome product to a target tissue, and